

08-18-00

A

Please type a plus sign (+) inside this box → ☐

Approved for use through 09/30/2000. OMB 0651-0032
Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

UTILITY PATENT APPLICATION TRANSMITTAL <small>(Only for new nonprovisional applications under 37 C.F.R. § 1.53(b))</small>	Attorney Docket No.	BEIERSDORF 634-WCG
	First Inventor or Application Identifier	Dr. Dieter WENNINGER
	Title	[See "Appendix"]
	Express Mail Label No.	EH976287555US

APPLICATION ELEMENTS See MPEP chapter 600 concerning utility patent application contents.	ADDRESS TO: Assistant Commissioner for Patents Box Patent Application Washington, DC 20231
---	---

1. <input checked="" type="checkbox"/> * Fee Transmittal Form (e.g., PTO/SB/17) (Submit an original and a duplicate for fee processing) 2. <input checked="" type="checkbox"/> Specification [Total Pages 38] (preferred arrangement set forth below) - Descriptive title of the invention - Cross References to Related Applications - Statement Regarding Fed sponsored R & D - Reference to Microfiche Appendix - Background of the invention - Brief Summary of the invention - Brief Description of the Drawings (if filed) - Detailed Description - Claim(s) - Abstract of the Disclosure 3. <input type="checkbox"/> Drawing(s) (35 U.S.C. 113) [Total Sheets] 4. Oath or Declaration [Total Pages 3] a. <input checked="" type="checkbox"/> Newly executed (original or copy) b. <input type="checkbox"/> Copy from a prior application (37 C.F.R. § 1.63(d)) (for continuation/divisional with Box 16 completed) i. <input type="checkbox"/> DELETION OF INVENTOR(S) Signed statement attached deleting inventor(s) named in the prior application, see 37 C.F.R. §§ 1.63(d)(2) and 1.33(b).	5. <input type="checkbox"/> Microfiche Computer Program (Appendix) 6. Nucleotide and/or Amino Acid Sequence Submission (if applicable, all necessary) a. <input type="checkbox"/> Computer Readable Copy b. <input type="checkbox"/> Paper Copy (identical to computer copy) c. <input type="checkbox"/> Statement verifying identity of above copies
---	--

ACCOMPANYING APPLICATION PARTS 7. <input checked="" type="checkbox"/> Assignment Papers (cover sheet & document(s)) 8. <input type="checkbox"/> 37 C.F.R. § 3.73(b) Statement <input checked="" type="checkbox"/> Power of Attorney (when there is an assignee) 9. <input type="checkbox"/> English Translation Document (if applicable) 10. <input checked="" type="checkbox"/> Information Disclosure Statement (IDS)/PTO-1449 <input checked="" type="checkbox"/> Copies of IDS Citations 11. <input checked="" type="checkbox"/> Preliminary Amendment 12. <input checked="" type="checkbox"/> Return Receipt Postcard (MPEP 503) (Should be specifically itemized) 13. <input type="checkbox"/> * Small Entity Statement(s) <input type="checkbox"/> Statement filed in prior application, Status still proper and desired (PTO/SB/09-12) 14. <input checked="" type="checkbox"/> Certified Copy of Priority Document(s) (if foreign priority is claimed) 15. <input checked="" type="checkbox"/> Other: Appendix, Form PTO-1449	
--	--

16. If a CONTINUING APPLICATION, check appropriate box, and supply the requisite information below and in a preliminary amendment:

☐ Continuation ☐ Divisional ☐ Continuation-in-part (CIP) of prior application No: _____

Prior application information: Examiner _____ Group / Art Unit: _____

For CONTINUATION or DIVISIONAL APPS only: The entire disclosure of the prior application, from which an oath or declaration is supplied under Box 4b, is considered a part of the disclosure of the accompanying continuation or divisional application and is hereby incorporated by reference. The incorporation can only be relied upon when a portion has been inadvertently omitted from the submitted application parts.

17. CORRESPONDENCE ADDRESS

☐ Customer Number or Bar Code Label (Insert Customer No. or Attach bar code label here) or ☒ Correspondence address below

Name	William C. Gerstenzang				
	NORRIS, McLAUGHLIN & MARCUS, P.A.				
Address	660 White Plains Road				
City	Tarrytown	State	New York	Zip Code	10591
Country	U.S.A.	Telephone	914-332-1700	Fax	914-332-1844

Name (Print/Type)	William C. Gerstenzang	Registration No. (Attorney/Agent)	27,552
Signature	<i>William C. Gerstenzang</i>	Date	Aug. 17, 2000

Burden Hour Statement: This form is estimated to take 0.2 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Box Patent Application, Washington, DC 20231

FEE TRANSMITTAL

for FY 2000

*Patent fees are subject to annual revision.
Small Entity payments must be supported by a small entity statement,
otherwise large entity fees must be paid. See Forms PTO/SB/09-12.
See 37 C.F.R. §§ 1.27 and 1.28.*

TOTAL AMOUNT OF PAYMENT (\$) 768.00

Complete if Known

Application Number	To Be Assigned
Filing Date	Herewith
First Named Inventor	Dr. Dieter WENNINGER
Examiner Name	To Be Assigned
Group / Art Unit	To Be Assigned
Attorney Docket No.	Beiersdorf 634-WCG

METHOD OF PAYMENT (check one)

1. ☒ The Commissioner is hereby authorized to charge indicated fees and credit any overpayments to:

Deposit Account Number 14-1263

Deposit Account Name NORRIS McLAUGHLIN & MARCUS, P.A.

- ☒ Charge Any Additional Fee Required
Under 37 CFR §§ 1.16 and 1.17

2. ☐ Payment Enclosed:
☐ Check ☐ Money Order ☐ Other

FEE CALCULATION

1. BASIC FILING FEE

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code (\$)	Fee Code (\$)	Fee Code (\$)	Fee Code (\$)		
101 690	201 345			Utility filing fee	690
106 310	206 155			Design filing fee	
107 480	207 240			Plant filing fee	
108 690	208 345			Reissue filing fee	
114 150	214 75			Provisional filing fee	

SUBTOTAL (1) (\$) 690.00

2. EXTRA CLAIM FEES

Total Claims		Extra Claims		Fee from below		Fee Paid	
14	-20** = 0	X	18	=	0		
4	-3** = 1	X	78	=	78		
Multiple Dependent			0	=	0		

**or number previously paid, if greater; For Reissues, see below

Large Entity		Small Entity		Fee Description
Fee Code (\$)	Fee Code (\$)	Fee Code (\$)	Fee Code (\$)	
103 18	203 9			Claims in excess of 20
102 78	202 39			Independent claims in excess of 3
104 260	204 130			Multiple dependent claim, if not paid
109 78	209 39			** Reissue independent claims over original patent
110 18	210 9			** Reissue claims in excess of 20 and over original patent

SUBTOTAL (2) (\$) 78.00

FEE CALCULATION (continued)


3. ADDITIONAL FEES

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code (\$)	Fee Code (\$)	Fee Code (\$)	Fee Code (\$)		
105 130	205 65			Surcharge - late filing fee or oath	
127 50	227 25			Surcharge - late provisional filing fee or cover sheet.	
139 130	139 130			Non-English specification	
147 2,520	147 2,520			For filing a request for reexamination	
112 920*	112 920*			Requesting publication of SIR prior to Examiner action	
113 1,840*	113 1,840*			Requesting publication of SIR after Examiner action	
115 110	215 55			Extension for reply within first month	
116 380	216 190			Extension for reply within second month	
117 870	217 435			Extension for reply within third month	
118 1,360	218 680			Extension for reply within fourth month	
128 1,850	228 925			Extension for reply within fifth month	
119 300	219 150			Notice of Appeal	
120 300	220 150			Filing a brief in support of an appeal	
121 260	221 130			Request for oral hearing	
138 1,510	138 1,510			Petition to institute a public use proceeding	
140 110	240 55			Petition to revive - unavoidable	
141 1,210	241 605			Petition to revive - unintentional	
142 1,210	242 605			Utility issue fee (or reissue)	
143 430	243 215			Design issue fee	
144 580	244 290			Plant issue fee	
122 130	122 130			Petitions to the Commissioner	
123 50	123 50			Petitions related to provisional applications	
126 240	126 240			Submission of Information Disclosure Stmt	
581 40	581 40			Recording each patent assignment per property (times number of properties)	
146 690	246 345			Filing a submission after final rejection (37 CFR § 1.129(a))	
149 690	249 345			For each additional invention to be examined (37 CFR § 1.129(b))	
Other fee (specify) _____					
Other fee (specify) _____					

* Reduced by Basic Filing Fee Paid

SUBTOTAL (3) (\$)

SUBMITTED BY

Name (Print/Type)	William C. Gerstenzang	Registration No. (Attorney/Agent)	27,552	Telephone	(914) 332-1700
Signature		Date	August 17, 2000		

WARNING:

Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

Burden Hour Statement: This form is estimated to take 0.2 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

Attorney Docket No. : Beiersdorf 634-WCG
: 6713-St-ar

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s) : Dr. Dieter WENNINGER, Dr. Werner KARMANN, Dr.
Gerhard BIR, Sven HANSEN, Heiko LEYDECKER, Stefanie
KAPROLAT, Ralf HIRSCH, Klaus MASSOW

For : ADHESIVE PACKAGING TAPE WITH NATURAL-RUBBER
HOT-MELT PRESSURE SENSITIVE ADHESIVE

Serial No. : To Be Assigned

Filed : Herewith

Art Unit : To Be Assigned

Examiner : To Be Assigned

August 17, 2000

BOX PATENT APPLICATIONS
Hon. Assistant Commissioner For Patents
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

In advance of prosecution, the Examiner is respectfully requested to amend the application as follows and consider the following remarks:

IN THE CLAIMS

Claim 1 (amended). Adhesive tape comprising a backing comprising an oriented thermoplastic film and a coating comprising a solventlessly prepared pressure-sensitive adhesive composition based on non-thermoplastic elastomers [such as natural rubber and tackifying resins, characterized in that] wherein the pressure-sensitive adhesive composition comprises a thermally labile crosslinking system.

Claim 2 (amended). Adhesive tape according to Claim 1, [characterized in that]

wherein the thermoplastic film comprises biaxially oriented HDPE, PVC or PET, monoaxially oriented polypropylene or biaxially oriented polypropylene.

Claim 3 (amended). Adhesive tape according to [Claims 1 and 2, characterized in that] Claim 1, wherein the adhesive composition comprises a mixture [comprising] of:

- | | |
|------------------------------|---|
| a) 100 parts by weight of | natural rubber |
| b) 70-120 parts by weight of | tackifying resins based on hydrocarbons |
| c) 5-30 parts by weight of | fillers |
| d) 2-20 parts by weight of | plasticizers |
| e) 0.1-15 parts by weight of | a crosslinker system |
| f) 0.5-5 parts by weight of | ageing inhibitors. |

Claim 4 (amended). Adhesive tape according to [one of Claims 1 to 3, characterized in that] Claim 1, wherein the crosslinker system is based on isocyanates, [such as diisocyanates or polyisocyanates,] and is used in [particular at] an amount of 0.1-5.0 parts by weight[, especially 0.1-2.0 parts by weight].

Claim 5 (amended). Adhesive tape according to [one of Claims 1 to 4, characterized in that] Claim 1, wherein the crosslinker system used comprises a mixture of 0.1-5 parts by weight of at least one photoinitiator, [used in particular at 0.1-5 parts by weight,] and 0.5 to 10 parts by weight of at least one polyfunctional (meth)acrylic ester[, used in particular at 0.5-10 parts by weight].

Claim 6 (amended). Adhesive tape according to [at least one of the preceding claims, characterized in that] Claim 1, wherein the pressure-sensitive adhesive composition is crosslinked by means of accelerated electrons or UV radiation.

Claim 7 (amended). Adhesive tape according to [at least one of the preceding claims, characterized in that] Claim 1, wherein a coat of a primer is applied between the thermoplastic film and the adhesive layer.

Claim 8 (amended). Adhesive tape according to [at least one of the preceding claims, characterized in that] Claim 1, wherein the thermoplastic film comprises a [coating, for example a] release coating.

Claim 9 (amended). Adhesive tape according to [at least one of the preceding claims, characterized in that] Claim 1, wherein the adhesion of the adhesive composition to the thermoplastic film is improved by means of corona treatment or[, preferably,] flame pretreatment.

Claim 10 (amended). A process for producing [an] the adhesive tape [according to at least one of the preceding claims, characterized in that] of Claim 1, wherein the pressure-sensitive adhesive composition based on non-thermoplastic elastomers is prepared continuously without solvent and without mastication in a continuously operating device having a filling section and a compounding section, the said process comprising

- a) feeding the solid components of the self-adhesive composition[, such as elastomers and resins,] into the filling section of the device, optionally feeding fillers, dyes, [and/or] crosslinkers, or combinations thereof,
- b) transferring the solid components of the self-adhesive composition from the filling section to the compounding section,
- c) adding the liquid components of the self-adhesive composition[, such as plasticizers, crosslinkers and/or further tackifying resins,] to the compounding section,
- d) preparing a homogeneous self-adhesive composition in the compounding section,

and

- e) discharging the self-adhesive composition.

Claim 11, please cancel.

Claim 12, please cancel.

Please add the following:

-- 13. The adhesive tape of Claim 1, wherein said adhesive composition comprises natural rubber and tackifying resins.

14. The adhesive tape of Claim 4, wherein said isocyanate is a diisocyanate or a polyisocyanate.

15. A method for sealing cartons which comprises sealing said cartons with an adhesive tape according to Claim 1 wherein said backing is based on biaxially oriented polyolefin, PVC or PET.

16. A method for bundling or palleting cardboard packaging wherein said cardboard packing is strapped with an adhesive tape according to Claim 1 wherein said backing is based on monoaxially oriented polefin. --

REMARKS

This Preliminary Amendment is being filed to eliminate multiple dependency and conform the claims to conventional format.


Favorable action is respectfully solicited.

ADDITIONAL FEE

Please charge any insufficiency of fees, or credit any excess, to Deposit Account No.
14-1263.

Respectfully submitted,

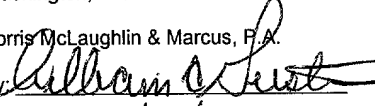
NORRIS, McLAUGHLIN & MARCUS, P.A.

By 
William C. Gerstenzang
Reg. No. 27,552

660 White Plains Road
Tarrytown, NY 10591-5144
(914) 332-1700

I hereby certify that this paper is being deposited with the
United States Postal Service as Express Mail, Label No.
EH976287555US, addressed to: BOX PATENT
APPLICATIONS, Hon. Assistant Commissioner of Patents,
Washington, D.C. 20231.

Norris McLaughlin & Marcus, P.A.

By 

Date: 8/17/00

09641014-031700

1. Personal Information	
Name	Mr. John Doe
Address	123 Main Street, Apt. 4B, New York, NY 10001
Phone	(212) 555-1234
Age	35
Gender	Male
Marital Status	Single
Education	Bachelor's Degree
Occupation	Software Engineer
Income	\$75,000 per year
Assets	Car, Savings, Stocks
Liabilities	Mortgage, Student Loans
Emergency Contact	John's Mother, (917) 555-5678
Signature	[Signature]
Date	10/26/2023
2. Financial Information	
Bank Name	Chase Bank
Account Type	Checking
Account Number	1234 5678 9010 1112
Routing Number	2222 3333 4444
Initial Deposit	\$1,000.00
Monthly Deposit	\$500.00
Monthly Withdrawal	\$200.00
Current Balance	\$1,300.00
Interest Rate	0.5%
Term	12 months
Payment Method	Automatic
Payment Date	1st of each month
Payment Amount	\$200.00
Signature	[Signature]
Date	10/26/2023
3. Additional Information	
Comments	I am interested in opening a new account and would like to know more about the services you offer.
Signature	[Signature]
Date	10/26/2023

Description

10

15

20

25

30

The unfavourable ageing behaviour and the profile of properties of such packaging tapes at increased temperatures, the poor thermal stability, result in premature opening of the boxes packed up using these packaging tapes, and/or a performance profile which is, in general, to some extent unfavourable for adhesive packaging tapes.

5

Various routes to the solvent-free preparation and processing of pressure-sensitive rubber adhesives are known.

10

All of the known processes are distinguished by very extensive rubber breakdown. For the further processing of the compositions to self-adhesive tapes, this requires extreme crosslinking conditions and also has the consequence of an application profile which is restricted to some extent, especially as regards the use of resultant self-adhesive tapes at relatively high temperatures.

15

The patent CA 698 518 describes a process for achieving production of a composition by adding high proportions of plasticizer and/or by simultaneously strong mastication of the rubber. Although this process can be used to obtain pressure-sensitive adhesives having an extremely high tack, user-compatible shear strength, even with a relatively high level of subsequent crosslinking, can be achieved only to a limited extent owing to the relatively high plasticizer content or else to the severe breakdown in molecular structure of the elastomer to a molecular weight average of $M_w \leq 1$ million.

20

25

The use of polymer blends, where besides non-thermoplastic natural rubber use is also made of block copolymers, in a ratio of approximately 1:1, is essentially an unsatisfactory compromise solution, since it results neither in high shear strengths when the self-adhesive tapes are used at relatively high temperatures nor in significant improvements relative to the properties described in the patent.

30

The deliberate industrial process of rubber breakdown under the combined action of shear stress, temperature and atmospheric oxygen is referred to in the technical literature as mastication and is generally carried out in the presence of chemical auxiliaries, which are known from the technical literature as masticating agents or peptizers, or, more rarely, as "chemical plasticizing aids".

In rubber technology, the mastication step is necessary in order to make it easier to integrate the additives.

According to Römpp (Römpp Lexikon Chemie – version 1.5, Stuttgart/New York: Georg Thieme Verlag 1998), mastication is a term used in rubber technology for the breakdown of long-chain rubber molecules in order to increase the plasticity and/or reduce the (Mooney) viscosity of rubbers. Mastication is accomplished by treating, in particular, natural rubber in compounders or between rolls at very low temperatures in the presence of mastication agents. The high mechanical forces which this entails lead to the rubber molecules being “torn apart”, with the formation of macroradicals, whose recombination is prevented by reaction with atmospheric oxygen. Mastication agents such as aromatic or heterocyclic mercaptans and/or their zinc salts or disulphides promote the formation of primary radicals and so accelerate the mastication process. Activators such as metal (iron, copper, cobalt) salts of tetraazaporphyrins or phthalocyanines permit a reduction in the mastication temperature. Mastication agents are used in the mastication of natural rubber in amounts of from about 0.1 to 0.5% by weight in the form of masterbatches which facilitate uniform distribution of this small amount of chemicals in the rubber composition.

Mastication must be clearly distinguished from the breakdown known as degradation which results in all of the standard solvent-free polymer technologies such as compounding, conveying and coating in the melt.

Degradation is a collective designation for various processes which alter the appearance and properties of plastics. Degradation may be caused, for example, by chemical, thermal, oxidative, mechanical or biological influences or else by exposure to radiation (such as (UV) light. Consequences are, for example, oxidation, chain cleavage, depolymerization, crosslinking, and/or elimination of side groups of the polymers. The stability of polymers to degradation may be increased by means of additives, for example by adding stabilizers such as antioxidants or light stabilizers.

Uncontrolled degradation often constitutes an unwanted phenomenon. It can be minimized by providing an inert gas atmosphere.

The use of non-thermoplastic elastomers is also described in JP 95 331 197, where use is made of an isocyanate-reactive natural rubber (polyisoprene grafted with maleic ester)

having an average molecular weight $M_w < 1$ million with aliphatic non-reactive hydrocarbon resins, which is crosslinked with blocked isocyanates (for example Desmodur CT); the mixture is initially crosslinked at 150°C for five minutes and following its subsequent coating onto PET film is cured at 180°C for several minutes (for example 15 minutes). This procedure clearly shows how complicated it is to achieve postcrosslinking if the natural rubber is subjected to excessive breakdown during the production process.

The patent application JP 95 278 509 discloses a self-adhesive tape in which the natural rubber is masticated to an average molecular weight of $M_w = 100,000$ to $500,000$ in order to obtain a coatable homogeneous mixture comprising hydrocarbon resins, rosin/rosin derivative resins and terpene resins, which can be processed readily at between 140°C and 200°C with a coating viscosity from 10 to 50×10^3 cps but requires an extremely high EBC dose (400 kGy) in order to ensure the shear strength necessary for its use.

For backing materials based on polyolefins, therefore, the system is unsuitable, since at the necessarily high beam doses there is significant backing deterioration. In the field of adhesive packaging tapes, such backing deterioration is unacceptable.

The term "adhesive packaging" or "adhesive tapes suitable for packaging" embraces on the one hand the adhesive packaging tapes for sealing cardboard packaging and on the other hand the strapping tapes used for firm bundling and palletization of cardboard packaging and other goods.

The use of exclusively non-thermoplastic rubbers as the elastomer component in the formulation of pressure-sensitive adhesives (PSAs) with the existing cost advantage possessed, for example, by natural rubbers over the standard commercial block copolymers, and with the outstanding properties, especially the shear strength of the natural rubber and corresponding synthetic rubbers, is also set out at length in the patents WO 94 11 175, WO 95 25 774, WO 97 07 963 and, correspondingly, US 5,539,033 and US 5,550,175.

In these cases, the additives customary in PSA technology, such as tackifier resins, plasticizers and fillers, are described.

The production process disclosed in each case is based on a twin-screw extruder which permits compounding to a homogenized PSA blend with the chosen process regime, involving mastication of the rubber and subsequent gradual addition of the individual additives with an appropriate temperature regime.

The mastication step of the rubber, which precedes the actual production process, is described at length. It is necessary and characteristic of the process chosen, since with the technology selected therein it is indispensable to the subsequent integration of the other components and to the extrudability of the blended composition. Also described is the feeding-in of atmospheric oxygen, as recommended by R. Brzoskowski, J.L. and B. Kalvani in *Kunststoffe* 80 (8), (1990), p. 922 ff., in order to accelerate rubber mastication.

This procedure makes it absolutely necessary to practice the subsequent step of electron beam crosslinking (EBC) and to use reactive substances as EBC promoters in order to achieve an effective crosslinking yield.

Both process steps are described in the abovementioned patents, but the EBC promoters selected also tend towards unwanted chemical crosslinking reactions at elevated temperatures. This limits the use of certain tackifying resins.

Owing to the unavoidable high production temperatures, compounding in a twin-screw extruder prevents the use of heat-activatable substances suitable for crosslinking the adhesive compositions, such as, for example, crosslinkers based on diisocyanate or other crosslinkers, since the chemical crosslinking reactions and secondary reactions that ensue in the process result in such a great increase in viscosity that the coatability of the resulting PSA composition is impaired and the composition therefore cannot be coated by means of nozzle coating.

The solvent-free hot-melt PSAs developed in recent years, based on non-thermoplastic elastomers, such as natural rubber or other high molecular mass rubbers, for example, therefore lack sufficient cohesion for the majority of applications, in the absence of a step of crosslinking the adhesive composition, and are therefore unsuited to use in the context of an adhesive packaging tape. The cause of this failure of non-crosslinked adhesive compositions based on natural rubber is the relatively large reduction in molecular weight as a result of processing, and/or as a result of the production process of the adhesive compositions based on natural rubber, and the resultant reduced or inadequate cohesion of the adhesive compositions.

The use of adhesive compositions of this kind based on natural-rubber hot-melt PSAs for adhesive tapes, especially adhesive packaging tapes, for sealing recycled-paper cardboard boxes or cardboard packaging, results in premature opening of the boxes. If there is sufficiently great tension across the lid of the box, caused by the pressure of the packaged material in the box or by the tension of the packaging material, which counters sealing, the adhesive tape becomes detached from the surface of the box and the box opens as a result of slippage of the adhesive packaging tape.

Not only for adhesive packaging tapes for sealing cardboard boxes, therefore, but also for other adhesive packaging tapes, such as strapping tapes, there is a need for sufficient cohesion of the adhesive composition based on natural rubber.

The cohesion and, with it, the packaging security afforded by adhesive tapes with an adhesive composition based on natural rubber can be improved either by crosslinking the rubber adhesive composition and/or by means of a preparation variant of the adhesive composition, in which the natural rubber used is broken down to a much-reduced extent and therefore has a higher molecular weight. This makes it possible to counter slippage of the adhesive tapes on the box surface, as described above.

Crosslinking possibilities for adhesive compositions based on natural rubber are known and are used to produce adhesive tapes. The crosslinking of the adhesive composition based on natural rubber may take place directly or by way of the use of a solvent-based primer which comprises a fraction of the suitable crosslinker.

Through migration of the crosslinking primer into the adhesive composition based on natural rubber, the adhesive composition can be durably crosslinked. Disadvantages of this known technology include the use of solvent-containing systems in the course of the coating process, the limiting pot lives of reactive primers of this kind, and the markedly increased expenditure in terms of occupational safety.

The crosslinking of the adhesive compositions based on natural rubber using primers proceeds relatively quickly, assisted by the dissolution effect of solvent-containing adhesive compositions.

The induced chemical crosslinking of an adhesive composition based on natural rubber by isocyanates, such as diisocyanates or polyisocyanates, for example, is known and is

used by various adhesive tape manufacturers in combination with solvent-containing adhesive compositions based on natural rubber. Adhesive compositions having improved cohesion, increased heat stability, and optimized packaging security are obtained by this means.

5

Owing to the unavoidably high process temperatures entailed in the use of known preparation processes for adhesive compositions based on natural rubber hot-melt PSAs, the use of chemical crosslinkers, such as isocyanate systems, for example, has to date been impossible owing to the crosslinking reactions which ensue.

10

A further possibility is the radiation crosslinking of the adhesive composition. Suitable for this purpose in principle are all types of ionizing radiation which can be used to produce, on the polymer molecules, free radicals or highly excited sites which subsequently react to give crosslinking sites. Particularly suitable are accelerated electrons having energies of from about 50 keV to 300 keV and ultraviolet (UV) rays having wavelengths of from about 200 to 400 nm, since they can be produced at high intensity and the penetration depth can be adapted to the coating thicknesses customary for adhesive tapes.

15

The fundamental possibilities of accelerated electrons for crosslinking PSA compositions are set out in US 2,956,904. One problem lies in the adverse effect on some substrates. In the case of OPP films, for instance, the light stability is greatly reduced, and PVC films undergo discoloration, especially after storage at elevated temperatures. Both can be reduced by means of specific stabilization, but this entails considerable additional cost.

20

UV radiation causes no substantial deterioration of the backing films. Crosslinking of the rubber adhesive composition requires the addition of at least one photoinitiator and one crosslinking promoter, as described in US 4,152,231. However, it is necessary to take account here of the absorption by fillers, colour pigments and other additives which leads to a decreasing degree of crosslinking in the depth of the coating. Weak layers of this kind may subsequently result in failure of the adhesive bond, especially on exposure to relatively high temperatures.

25

30

Monoaxially and biaxially oriented films based on polypropylene are used in large amounts for adhesive packaging tapes. Whereas biaxially oriented films based on polypropylene are preferred for the use of an adhesive packaging tape for carton sealing,

35

monoaxially oriented polypropylene films find application in the strapping tapes segment. These films are notable for very high tensile strength and low extension in the longitudinal direction and are highly suited to bundling and to the holding-together of pallets. When using adhesive compositions which exhibit inadequate cohesion or inadequate anchoring to the film, the slippage of the strapping tapes results in slipping of the pallets and thus in inadequate securement of the pallets. Insufficient anchoring of the adhesive composition to film leads, when using carton sealing tapes, to premature opening of the packed-up boxes.

- 10 A natural rubber hot-melt pressure-sensitive adhesive prepared without solvent, and thus in an environmentally gentle way, in combination with a backing film made from environmentally gentle polypropylene, suggests on conceptual grounds the desire for further environmentally compatible product and production technologies – in so far as they become additionally necessary. Polypropylene, as a nonpolar, saturated hydrocarbon, is not immediately suitable for providing sufficiently firm anchoring of a primer or a natural rubber adhesive composition. This can be remedied to a certain extent by activating oxidative pretreatment of the backing – for example, by corona treatment or flame treatment. Both methods are environmentally gentle, by saving on raw materials, and are attracting very great interest as cost-reducing alternatives to wet-chemical primer coating. The use of a primer alone generally does not solve the anchoring problem on polypropylene; the primer itself requires activation of the backing for its fixing on this substrate. The great efficiency of flame treatment, especially at high web speeds, has been known for a long time, although it is only since modern, readily controllable flaming units have become available that flaming has emerged, in its industrial importance, from the shadow of corona treatment. If the anchoring strength requirements are not too great, flame treatment does in certain cases make it possible to omit the use of a primer. The realization of anchoring, in a manner which meets the requirements, by means of backing flame treatment alone, in the specific case of a natural-rubber hot-melt pressure-sensitive adhesive on polypropylene films for high-quality adhesive packaging tapes, is in ideal harmony with the environmental protection aspects outlined.

Thermoplastic films based on polyethylene terephthalate (PET) or polyvinyl chloride (PVC) are likewise known and are used by various manufacturers to produce adhesive tapes. In this context, films based on PET, in particular, are distinguished by high

elongation at break and thermal stability of from 130°C to 175°C, and resistance to dilute alkalis and acids. Moreover, films based on polyesters possess very high abrasion resistance and penetration resistance, but are less widespread in the field of adhesive packaging tapes owing to the relatively high price in relation to polyolefin-based films.

5

The PVC may be obtained by emulsion, suspension or bulk polymerization. Copolymers based on PVC/vinyl acetate are also known.

10

Films based on PVC in combination with solvent-containing adhesive compositions based on natural rubber are offered as adhesive packaging tapes by various manufacturers. In this case, owing to the better thermal stability, unplasticized PVC films are used which have a good thermal stability of up to max. 105°C.

15

In the production of adhesive tapes with adhesive compositions based on natural rubber, both water-based and solvent-containing primers are used as adhesion promoters between adhesive composition and backing film. These adhesion promoters that are used possess in part a crosslinking effect on the natural-rubber-based adhesive composition which is applied from solution.

20

Furthermore, the films based on unplasticized PVC are distinguished by moderate penetration strength and a partial resistance to petrol, oil and alcohols. In general, the anchoring of primers to PVC films without corona treatment is better than with other thermoplastic films, such as polyolefin-based films, for example. The reason for this is the high surface energy even without pretreatment [approximately 39 dyn/cm] and the rough surface structure of PVC films. For sufficient anchoring of adhesive compositions based on natural rubber, a primer-coated PVC film is advisable.

25

30

It is an object of the present invention to provide adhesive tapes comprising films based on oriented thermoplastic films and solventlessly prepared adhesive compositions based on natural rubber. Furthermore, the new adhesive tapes should unroll readily and have a good bond between adhesive composition and backing film. Furthermore, the new adhesive tapes should be suitable either as adhesive packaging tapes or as strapping tapes for standard commercial cardboard packaging.

This object is achieved by means of an adhesive tape as set out in the main claim. The subsidiary claims relate to advantageous developments of the subject-matter of the invention. Furthermore, the invention provides processes for producing and proposals for using the adhesive tape of the invention.

5

The invention accordingly provides an adhesive tape comprising a backing comprising an oriented thermoplastic film and a coating comprising a solventlessly prepared pressure-sensitive adhesive composition based on non-thermoplastic elastomers such as natural rubber and tackifying resins, the pressure-sensitive adhesive composition comprising a thermally labile crosslinking system.

Films based on polyesters, for example polyethylene terephthalate, are likewise known and may also be used to produce the adhesive tapes of the invention. The thicknesses of the films based on PET are between 20 and 100 μm , in particular between 30 and 50 μm .

Films based on oriented polyolefins are known and belong to the prior art. Monoaxially and biaxially oriented films based on polyolefins are used in large amounts for adhesive packaging tapes, strapping tapes, and other adhesive tapes. Films based on oriented polyethylene or oriented copolymers comprising ethylene and/or polypropylene units are also known. All of these films may be used as backing film in accordance with the invention.

Monoaxially oriented polypropylene is noted for its very high tensile strength and low elongation in the longitudinal direction and is used, for example, to produce strapping tapes. Monoaxially oriented films based on polypropylene are preferred for producing the adhesive tapes of the invention for bundling and palletizing cardboard packaging and other goods. The thicknesses of the monoaxially oriented films based on polypropylene are preferably between 25 and 200 μm , in particular between 40 and 130 μm .

Films comprising monoaxially oriented polypropylene are particularly suitable for producing the adhesive tapes of the invention. Monoaxially oriented films are predominantly single-layered, although multilayer monoaxially oriented films may also be produced in principle. The known films are predominantly one-, two- and three-layer films, although the number of layers chosen may also be greater.

For the production of the adhesive tapes of the invention for secure carton sealing, preference is given to biaxially oriented films based on polypropylene with a draw ratio in the longitudinal (machine) direction of between 1:4 and 1:9, preferably between 1:4.8 and 1:6, and a draw ratio in the transverse (cross) direction of between 1:4 and 1:9, preferably between 1:4.8 and 1:8.5.

The moduli of elasticity achieved in the longitudinal direction, measured at 10% elongation in accordance with ASTM D882, are usually between 1000 and 4000 N/mm², preferably between 1500 and 3000 N/mm².

The thicknesses of the biaxially oriented films based on polypropylene are in particular between 15 and 100 µm, preferably between 20 and 50 µm.

Biaxially oriented films based on polypropylene may be produced by means of blown film extrusion or by means of customary flat film units. Biaxially oriented films are produced both with one layer and with a plurality of layers. In the case of the multilayer films, the thickness and composition of the different layers may also be the same, although different thicknesses and compositions are known.

Particularly preferred for the adhesive tapes of the invention are single-layer, biaxially or monoaxially oriented films and multilayer biaxial or monoaxial films based on polypropylene which have a sufficiently firm bond between the layers, since delamination of the layers in the course of the application is disadvantageous.

Films based on unplasticized PVC are likewise known and are used by various adhesive tape manufacturers to produce adhesive packaging tapes. Plasticized PVC, owing to the inadequate thermal stability of max. 60°C, is unsuited to the production of adhesive packaging tapes but may, if desired, also be used.

For the adhesive tapes of the invention for secure carton sealing it is preferred to use films based on unplasticized PVC. The thicknesses of the films are preferably between 30 and 100 µm, in particular between 35 and 50 µm. The adhesive tapes of the invention are used as packaging tapes for carton sealing.

In one preferred embodiment of the adhesive tape, the adhesion of the adhesive composition on the thermoplastic film based on polyolefins is improved by means of

corona treatment or, especially, by means of flame pretreatment, since, especially, the surfaces of films based on oriented polyolefins may be treated by means of these widely known processes, such as corona treatment or flame treatment. Preference is given to surface treatments by flame pretreatment. An overview of the processes for surface treatment is contained, for example, in the article "Surface pretreatment of plastics for adhesive bonding"/A. Kruse; G. Krüger, A. Baalmann and O. D. Hennemann; J. Adhesion Sci. Technol., Vol. 9, No. 12, pp. 1611-1621 (1995).

The biaxially or monoaxially oriented films based on polyolefins for the adhesive tapes of the invention are preferably corona- and/or flame-pretreated on the side facing the adhesive composition in order to obtain sufficient anchoring of the adhesive composition on the film. Flame pretreatment is the preferred form of film pretreatment for films based on oriented polyolefins.

The flaming installation comprises at its core a coolable burner unit and a likewise coolable treatment roll, which serves to guide the web. While the burner temperature in the course of flaming is held at between 30°C and 40°C, the cooling of the treatment roll ensures temperatures there of between 10°C and 15°C. The film web is guided and treated between burner and treatment roll under defined conditions. The flaming of the abovementioned polyolefin films takes place by way of a fuel gas/air mixture in optimized composition. The fuel gas fraction comprises gaseous hydrocarbons such as propane or butane or hydrocarbon mixtures in the form, for example, of standard commercial natural gas. Depending on the precise composition of the fuel gas/air mixture, flame temperatures of between 750°C and 900°C are achieved. The distance of the burner unit from the treatment roll may be regulated between 2 mm and 7.5 mm. Web speeds can be set within the limits from 30 m/min up to several hundred m/min.

An upper limit results from the capacity of the available installation, i.e. the total installation including coating section in the case of in-line backing treatment, or the flaming installation in the case of off-line backing pretreatment.

The lower limit is governed by the thermal sensitivity of the film material. The energy input can be varied by way of the volume flow of the mixture, at given settings for mixture composition, burner distance and web speed, within limits from 20 m³/h to 50 m³/h. In addition, the effective flaming width can be adjusted to different film widths by way of adjustable burner boundary.

Backing material flamed under optimized parameters is coated with natural-rubber hot-melt pressure-sensitive adhesive no later than one day following pretreatment. A similar procedure is followed for comparative material which has been appropriately corona-pretreated. With suitable plant construction, in-line flaming or corona pretreatment prior to coating of the adhesive composition based on natural-rubber hot-melt pressure-sensitive adhesives is possible and leads to a further improvement in the anchoring of the composition.

The anchoring strength of the adhesive composition is tested on finished adhesive tapes, especially after a matured state has been reached. In the course of this testing, the question as to on-specification anchoring strength is of just as much interest as the comparison between flaming and corona pretreatment.

It is found that flaming of the backing brings about far better anchoring of natural-rubber hot-melt pressure-sensitive adhesives than a corona pretreatment appropriate for comparison. Anchoring on the pretreated backing undergoes further, considerable increase if the natural-rubber hot-melt pressure-sensitive adhesive comprises isocyanate crosslinkers. Especially in combination with the flaming of the backing, it is in some cases possible by this means to forego the use of a primer, which in turn permits the omission of a drying tunnel in the coating process, this process thus being totally solvent-free.

The adhesive composition itself preferably comprises a mixture comprising:

- | | |
|------------------------------|---|
| a) 100 parts by weight of | natural rubber (granulated) |
| b) 70-120 parts by weight of | tackifying resins based on hydrocarbons |
| c) 5-30 parts by weight of | fillers, for example chalk and/or organic or inorganic pigments |
| d) 2-20 parts by weight of | plasticizers, for example mineral oil |
| e) 0.1-15 parts by weight of | a crosslinker system |
| f) 0.5-5 parts by weight of | ageing inhibitors. |

If desired, the mixture may be further supplemented by

- | | |
|----------------------------|---------------------------------|
| g) 0-10 parts by weight of | organic and inorganic pigments. |
|----------------------------|---------------------------------|

The application rate of the adhesive composition to the backing material, especially a thermoplastic film made from oriented polyolefins, PVC or PET, is preferably from 10 to 50 g/m². In a further preferred embodiment, the layer application established is from 14 to 25 g/m².

5

The adhesive tapes of the invention may comprise further layers. The use of appropriate primers, disposed between adhesive layer and film, is possible provided it does not reduce the good anchoring, in accordance with the invention, between the adhesive composition based on natural-rubber hot-melt pressure-sensitive adhesive and the flame- or corona-treated film. The use of an appropriate coating, for example a release coating, is possible.

10

15

Adhesive compositions based on natural rubber have been known for a long time and are widely used for adhesive packaging tapes. An overview of the adhesive compositions and their use in the field of pressure sensitive adhesives can be found in "Natural Rubber Adhesives" (G. L. Butler in Handbook of Pressure Sensitive Adhesive Technology, Third Edition, edited by Donatas Satas, Van Nostrand Reinhold New York, pp.261-287).

20

The high molecular weight and the unsaturated molecule chains of the natural rubber govern its advantages in terms of material properties and at the same time the problems associated with its processing. For performance-capable adhesive tapes, natural rubber can to date be processed only by solvent technology. This technology has limits in economics, environmental acceptance, and the further development of the adhesive compositions for adhesive packaging tapes. Known solvent-free technologies, such as, for example, the use of thermoplastic block copolymers comprising polystyrene blocks, are based on low melt viscosities, which lead to poor thermal stability. These low melt viscosities are not present in the case of natural rubber. Furthermore, high process temperatures in the case of natural rubber result in thermooxidative damage, owing to the unsaturated natural rubber molecules. The combination of planetary roll extruders as a suitable compounder unit and the coating of the adhesive composition by means of a roll applicator permits the use of solventlessly prepared adhesive compositions based on natural rubber for adhesive packaging tapes and combines the efficiency of the hot-melt

25

30

pressure-sensitive adhesive technology with the advantages of the natural rubber technology.

5 The adhesive compositions of the invention based on natural rubber are prepared and applied without solvent and comprise in particular crepe, SSR or ADS, TSR grades of natural rubber as the natural rubber component. Particular preference is given to ADS and TSR rubber grades as the natural rubber component.

10 In order to be able to incorporate different components into the natural rubber fraction of the adhesive composition, to date the drastic breakdown of the molecular weight of the rubber has been absolutely necessary for producing the adhesive composition. This process (mastication) takes place by way of special compounders and is particularly shear-intensive. The extent of the breakdown in molecular weight can be controlled within certain limits via the input of shear energy, temperature, and controlled action of oxygen.

15 The use of a planetary roll extruder for producing the adhesive compositions based on natural rubber makes it possible to produce homogenized adhesive compositions by means of very high shear rates in conjunction with excellent heat exchange and a very short residence time of the composition. This brings about a relatively low reduction in the molecular weight of the natural rubber at simultaneously low process temperatures of
20 from 90 to 110°C.

25 Tackifying resins based on hydrocarbons are known and are in very widespread use for adhesive compositions for adhesive packaging tapes. A description and possible uses in connection with natural-rubber-based adhesive compositions is described, for example, in "Natural Rubber Adhesives" (G. L. Butler in Handbook of Pressure Sensitive Adhesive Technology, Third Edition, edited by Donatas Satas, Van Nostrand Reinhold New York, pp.263-266). Hydrocarbon-based tackifying resins are available commercially with a range of different properties such as, for example, different chemical structure, colour
30 grades and softening points.

For the solvent-free, natural-rubber-based adhesive compositions of the invention, hydrocarbon-based tackifying resins are used. The solvent-free adhesive composition based on natural rubber comprises from 70 to 120 parts of tackifying resin and preferably from 80 to 110 parts of tackifying resin per 100 parts of natural rubber.

The use of tackifying resins based on mixtures of different hydrocarbon resins and on mixtures of hydrocarbon resins of the invention with other known resins is also in accordance with the invention provided that it does not alter the characteristic profile of properties of the adhesive tapes of the invention in a manner detrimental to those properties. Possible, for example, are mixtures of hydrocarbon resins with fairly small amounts of tackifying resins based on rosin or modified rosin or on phenolic resins, other natural resins, resin esters or resin acids.

The use of plasticizers as additives to adhesive compositions for adhesive packaging tapes is known.

For the adhesive tapes of the invention, use is made of from 2 to 20 parts of plasticizer, based on paraffinic and naphthenic oils, oligomers such as oligobutadienes and oligoisoprenes, liquid nitrile rubbers, liquid terpene resins, vegetable and animal oils and fats and phthalates, per 100 parts of natural rubber.

Particularly preferred for the adhesives of the invention are solvent-free adhesive compositions based on natural rubber and comprising from 4 to 10 parts of plasticizer. Particular preference is given to the use of paraffinic and naphthenic oils.

Ageing inhibitors for adhesive compositions based on natural rubber are known. Three different kinds of ageing inhibitor are used in particular as antioxidants for adhesive compositions: ageing inhibitors based on amines, on dithiocarbamates, and on phenols.

Ageing inhibitors based on phenols are very effective under the influence of UV radiation and sunlight.

For the adhesive tapes of the invention, ageing inhibitors based on phenols are used. The adhesive composition comprises in particular, from 0.5 to 5 parts of an appropriate ageing inhibitor, based on phenols, per 100 parts of natural rubber. Also in accordance with the invention is the use of other types of ageing inhibitor, such as, for example, ageing inhibitors based on amines and dithiocarbamates.

Organic and inorganic pigments for adhesive compositions based on natural rubber are known. For the colouring of adhesive compositions based on natural rubber, use is made in particular of titanium dioxide or titanium dioxide in combination with different-coloured colour pigments.

- 5 For the adhesive tapes of the invention, suitable organic and/or inorganic colour pigments are used. The adhesive composition comprises from 0 to 10 parts of an organic and/or inorganic colour pigment, per 100 parts of natural rubber. Particular preference is given to the use of from 0 to 7 parts of an appropriate organic or inorganic colour pigment per 100 parts of natural rubber. Also in accordance with the invention is the use of pigments
10 based on mixtures of different inorganic and organic pigments.

- Suitable fillers for adhesive compositions based on natural rubber are known. In this instance, calcium carbonate (chalk), titanium dioxide, zinc oxide, clay or pigments are
15 used in particular. For the adhesive tapes of the invention, fillers based on chalk, titanium dioxide, clay or pigments are used. The adhesive composition comprises from 10 to 30 parts of fillers, in particular from 10 to 20 parts of fillers, per 100 parts of natural rubber. Particular preference is given to the use of calcium carbonate (chalk) as filler. Also in accordance with the invention is the use of fillers based on mixtures of different
20 fillers.

- The crosslinking of adhesive compositions based on natural rubber is known and is described, for example, in "Natural Rubber Adhesives" G. L. Butler in Handbook of
25 Pressure Sensitive Adhesive Technology, Third Edition, edited by Donatas Satas, Van Nostrand Reinhold New York, pp.269-276. The adhesive composition based on the natural rubber is crosslinked either by using chemical crosslinkers or by using physical crosslinking methods such as electron beam crosslinking or crosslinking by means of UV radiation.

- 30 For the purpose of chemically-thermally induced crosslinking it is possible in general to use known crosslinkers such as accelerated sulphur systems or sulphur donor systems, isocyanate systems, reactive melamine resins, formaldehyde resins and (optionally halogenated) phenol-formaldehyde resins and/or reactive phenolic resin crosslinking

systems or diisocyanate crosslinking systems with the appropriate activators, epoxidized polyester resins and acrylate resins, and combinations thereof.

Crosslinkers based on isocyanates, such as diisocyanate systems or polyisocyanates, for example, are particularly preferred. The use of these crosslinking systems is known in principle and is described, for example, in GB 1,234,860.

The adhesive compositions of the adhesive tapes of the invention comprise from 0.1 to 15 parts of a suitable crosslinker per 100 parts of natural rubber.

In one preferred embodiment the adhesive composition comprises from 0.1 to 5 parts of an isocyanate, such as diisocyanates or polyisocyanates, in a particularly preferred embodiment from 0.5 to 2.0 parts of isocyanates, per 100 parts of natural rubber. The amount of suitable crosslinker may be adapted to the amount of tackifying resin according to the invention.

Radiation crosslinking has the advantage of very rapidly achieving a high degree of crosslinking which then remains stable over a long period. Consequently, in-line crosslinking is possible even at high coating speeds of several 100 m/min, and in contrast to chemical crosslinking the quality of the adhesive properties can be checked directly after the end of production.

By controlling the intensity of radiation it is possible to establish the desired degree of crosslinking within wide ranges, independently of the web speed.

Rubber adhesive compositions may be crosslinked by means of accelerated electrons without specific additives. In order to raise the crosslinking yield it is possible to add crosslinking promoters such as polyfunctional (meth)acrylates, thiols or bismalimides.

The available installations and the most important process parameters have been disclosed elsewhere (W. Karmann, J. of Industrial Irradiation Technology 1(4) (1983) pp. 305-323).

Owing to the damage caused to numerous important packaging tape backings such as OPP and PVC films by EBC, the crosslinking of the adhesive composition by UV radiation is particularly advantageous. For this purpose it is necessary to add photoinitiators and crosslinking promoters to the rubber adhesive composition. Examples of photoinitiators which may be used include aromatic ketones, benzoin and derivatives, benzil and dialkyl

ketals derived therefrom, thioxanthans, anthraquinones, anthracenes and phosphine oxides. Suitable crosslinking promoters include acrylic and methacrylic esters of polyfunctional alcohols such as hexanediol, trimethylolpropane and the ethoxylated and dimerized derivatives thereof, pentaerythritol and the dimer, polyethylene glycol and polypropylene glycol. For hot-melt pressure-sensitive adhesives, compounds with relatively high molecular weights are generally preferred owing to their lower vapour pressure. Acrylic esters usually give higher reaction rates.

In the case of UV crosslinking, it is necessary to bear in mind the absorption of the radiation by the photoinitiator and by organic and inorganic pigments in the adhesive composition, which leads to a sharp decrease in the degree of crosslinking with penetration in the adhesive composition layer. Excessive crosslinking on the surface of the adhesive composition has the consequence of unfavourable adhesive properties.

This can be avoided, or at least reduced, if the adhesive composition is irradiated through the backing film or from both sides. The effect may also be reduced by reducing the shortwave component of the radiation with a wavelength of less than about 300 nm by means of various measures. Suitable radiation tubes for this purpose are available on the market. The component may also be eliminated by means of appropriate filtering.

The radiation intensity can be controlled by incorporating radiation tubes and also, within certain limits, by controlling the output of the radiation emitters, as described by A. Beying, RadTech Europe '97.

Advantageously, moreover, the effect of atmospheric oxygen on the surface of the adhesive composition to be crosslinked is excluded. This can be achieved by blanketing with inert gases such as nitrogen, carbon dioxide and/or noble gases such as argon or by covering with a release liner (release paper or release films) which may, if desired, be transparent.

In one further embodiment, accordingly, the adhesive compositions of the adhesive tapes of the invention comprise not only from 0.1 to 5 parts, preferably from 0.5 to 3 parts, of at least one photoinitiator but also from 0.5 to 10 parts, preferably from 1 to 5 parts, of a polyfunctional (meth)acrylic ester. The amount of appropriate crosslinker may be adapted to the amount of the resin that is used.

For the electron beam crosslinking of adhesive compositions with coatings of from 10 to 50 g/m², it is preferred to use lead-shielded single-stage electron accelerators with voltages of from about 50 to 300 kV, which owing to their relatively small dimensions are

easy to integrate into the coating installations. The types available on the market are described by G.G. Skelthorne ("Electron Beam Processing" in Vol. 1 "Chemistry & Technology of UV & EB Formulations for Coatings, Inks & Paints" publ. Sita Technology, London 1991).

- 5 UV radiation installations comprise one or more gas discharge tubes, which are mostly provided with elliptical or parabolic reflectors and with an effective cooling means. An overview of the different tubes and the other components is given by R.E. Knight ("UV Curing Equipment and Applications" in Vol. 1 "Chemistry & Technology of UV & EB Formulations for Coatings, Inks & Paints" publ. Sita Technology, London 1991).

10

The production of the adhesive composition based on natural rubber of the adhesive tapes suitable for packaging may take place by means of a single-layered or multilayered production. Suitable in this context are, for instance, production in the batch process and
15 the use of twin-screw extruders. Overall it is possible to coat adhesive compositions of this kind onto thermoplastic films by means of the abovementioned roll applicator.

15

In this case, in the context of batchwise production, a pre-batch is prepared using a Banbury compounder and then mixing is completed with the aid of a kneading machine
20 after the complete raw-material components have been added.

20

The invention then embraces a process for continuous solvent-free and mastication-free preparation of pressure-sensitive adhesive compositions based on non-thermoplastic elastomers in a continuously operating device having a filling section and a compounding
25 section, comprising the following steps:

25

- a) feeding the solid components of the self-adhesive composition, such as elastomers and resins, into the filling section of the device, optionally feeding fillers, dyes and/or crosslinkers,
- 30 b) transferring the solid components of the self-adhesive composition from the filling section to the compounding section,
- c) adding the liquid components of the self-adhesive composition, such as plasticizers, crosslinkers and/or further tackifying resins, if desired in the melted state, to the compounding section,
- 35 d) preparing a homogeneous self-adhesive composition in the compounding section,

- e) discharging the self-adhesive composition, and
- f) coating the self-adhesive composition onto a material in web form, the coating of the web-form material being carried out with a multi-roll applicator, in particular with a 2- to 5-roll applicator, especially with a 4-roll applicator, so that the self-adhesive composition is shaped to the desired thickness as it passes through one or more roll nips, it being possible to set the rolls of the applicator individually to temperatures of from 20°C to 150°C.

The use of a planetary roll extruder as a continuously operating device has been found to be particularly advantageous, the compounding section of the said device comprising preferably at least two, but with particular preference three, coupled roll cylinders, it being possible for each roll cylinder to have one or more separate temperature control circuits.

Unlike otherwise conventional production processes, in the planetary roll extruder in accordance with the process of the present invention, in particular, there is no property-impairing mastication of the non-thermoplastic elastomers, since in this case they are not subjected separately to the effect of high shear energy but instead are always processed together with one or more liquid components. These liquid components may include plasticizers such as oils, for example, and also resins which melt only during the compounding process under the action of shear energy and/or external heating. The presence of these liquid components limits the extent of frictional energy such that it is possible to avoid the mastication of the rubber, i.e. the molecular weight breakdown of the elastomers, and the high resultant compounding temperatures.

Furthermore, planetary roll extruders have extremely large areas where heat exchange, material exchange and surface renewal take place, by which means it is possible to dissipate rapidly the frictional shear energy and thus to avoid undesirably high product temperatures.

Planetary roll extruders have been known for a fairly long time and were first used in the processing of thermoplastics such as PVC, for example, where they were used primarily to supply the downstream units such as, for example, calenders or roll mills. As a

consequence of their advantage of the great renewal of surface area for material exchange and heat exchange, by means of which the frictional energy can be dissipated rapidly and effectively, and because of the low residence time and the narrow residence-time spectrum, their use in recent times has been extended, inter alia, to compounding processes which require a particularly temperature-controlled regime.

Depending on manufacturer, planetary roll extruders are available in various designs and sizes. The diameters of the roll cylinders, depending on the desired throughput, are typically between 70 mm and 400 mm.

10 Planetary roll extruders generally have a filling section and a compounding section.

The filling section comprises a conveying screw to which all of the solid components are fed continuously. The conveying screw then passes the material to the compounding section. The region of the filling section, together with the screw, is preferably cooled in order to prevent baking-on of material on the screw. Alternatively, there are designs without a screw section, where the material is fed directly between central spindles and planetary spindles. However, this is not significant for the effectiveness of the process of the invention.

20 The compounding section comprises a driven central spindle and a plurality of planetary spindles which rotate around the central spindle within a roll cylinder with internal helical gearing. The rotary speed of the central spindle, and hence the rotational speed of the planetary spindles, can be varied and is therefore an important parameter for controlling the compounding process.

25 The materials are circulated between the central and planetary spindles, and between the planetary spindles and the helical gearing of the roll section, so that under the effect of shear energy and external heating the materials are dispersed to form a homogeneous compound.

30 In this context, reference is made to the patent applications and, respectively, utility model DE 196 31 182, DE 94 21 955, DE 195 34 813, DE 195 18 255, DE 44 33 487, which offer an overview of the state of the art in the field of planetary roll extruders.

35 It is advantageous to use a planetary roll extruder whose compounding section has been extended by coupling at least two roll cylinders. Firstly, despite the presence of friction-reducing components for the purpose of avoiding mastication of the rubber, the complete

digestion of the elastomer components, and the desired homogenizing and dispersing performance at economic throughput rates, is possible by this means; secondly, the coupling of, preferably, separately temperature-controlled roll cylinders permits a balanced temperature regime in the process, so allowing the use of heat-activatable crosslinker systems.

Whereas in the front compounding section of the planetary roll extruder the roll cylinders are advantageously heated at temperatures above the melting point of the resins used, the rear compounding section is advantageously cooled in order to reduce the product temperature. By this means, the residence time of the self-adhesive composition at relatively high temperatures is kept as short as possible, thereby preventing activation of the thermal crosslinker systems present in the self-adhesive composition.

In the first step of the process, a composition comprising the elastomers and the known additives required for the preparation of self-adhesive compositions, such as fillers, ageing inhibitors, plasticizers and tackifying resins, is prepared without solvent in a planetary roll extruder, the composition having a final temperature of less than 150°C, preferably less than 130°C, with very particular preference between 70°C and 110°C. The overall residence time of the composition within the planetary roll extruder should not exceed a level of three minutes. The resulting hot-melt adhesive composition has a viscosity of between 300 and 1500 Pa*s in particular a viscosity of between 800 and 1200 Pa*s at 130°C and a shear rate of 100 rad/s.

In the second step of the process, which advantageously takes place in conjunction with the compounding step in the planetary roll extruder, the pressure-sensitive hot-melt adhesive composition prepared in accordance with the invention is coated without solvent onto a backing in web form, using an applicator unit.

In accordance with the invention it is further proposed to carry out the coating of the web-form material without solvent using a roll coating applicator unit or muliti-roll coating calenders comprising preferably three, with particular preference four, coating rolls, the self-adhesive composition being shaped to the desired thickness as it passes through one or more roll nips before transfer to the web-form material. This coating process is particularly preferable when the viscosities of the self-adhesive composition exceed levels

of 5000 Pa*s at a shear rate of 1 rad/s, since in this case coating with extrusion dies no longer provides the necessary accuracy in terms of the amount of composition applied.

Depending on the nature of the web-form backing material to be coated, coating may take place in a co-rotating or counter-rotating process.

5

Coating on roll coating applicator units or multiroll coating calenders is possible at temperatures below 100°C, so that even self-adhesive compositions containing heat-activatable crosslinkers can be coated. The roll which carries the backing is cooled at from 5°C to 25°C, preferably from 10 to 15°C. It is possible as a result to coat heat-sensitive backings, such as unplasticized PVC for example.

10

The rolls of the applicator unit may be set individually at temperatures from 20°C to 150°C.

15

The preferred 4-roll applicator unit is formed of a metering roll, a knife-coating roll, which determines the thickness of the layer on the backing material and is arranged parallel to the metering roll, and a transfer roll, which is situated below the metering roll. On the placement roll, which together with the transfer roll forms a second roll nip, the composition and the web-form material are brought together. The placement roll is cooled at a temperature of from 5 to 25°C, preferably from 10 to 15°C.

20

It is thus possible, without wishing hereby to restrict the concept of the invention, for the differential speeds in the case of an appropriate 4-roll applicator unit to be as follows:

25

Roll	Differential speed with respect to the relative web speed [%]
Metering roll	from 0.1 to 25 in particular from 1 to 2
Knife-coating roll	from 10 to 100 in particular from 30 to 40
Transfer roll	from 10 to 100 in particular from 75 to 90

For the purpose of increased freedom of the coated adhesive composition from gas bubbles, it is possible to install a vacuum devolatilizer between planetary roll extruder and applicator unit, for example a vacuum chamber, a devolatilizing extruder or the like.

5

The adhesive tapes of the invention are suitable for a large number of adhesive tape applications. One important field of application is that of packaging applications. The adhesive tapes are suitable for use as carton sealing tapes, general adhesive packaging tapes, strapping tapes, and adhesive tapes for sealing plastic packaging and plastic bags. The adhesive tapes are suitable for pallet securement. Further applications are the bundling of loose goods and goods for transportation, such as pipes, wooden strips, etc., for example. The adhesive tapes of the invention may be used to secure, for example, refrigerators and other electrical and electronic appliances when in transit. Further applications are label protection, surface protection, in the construction sector, for example, and tear-open strips for packaging. Applications within the office sector are also possible.

10

15

20

The intention of the text below is to illustrate the invention, with reference to examples, without wishing unnecessarily to restrict the invention.

25

Examples

Example a: Film

A biaxially oriented film based on polypropylene is used, from Radici.

30

Film thickness	25 μm
Designation	Radil T 25 NT (NT = Not treated)
Company/manufacturer	Radici

Elongation at break, longitudinal	130%
Elongation at break, transverse	50%
Modulus of elasticity longitudinal	2500 N/mm ²
Modulus of elasticity transverse	4000 N/mm ²
Thermal shrinkage at 130°C for 60 s.	longitudinal <5%
Thermal shrinkage at 130°C for 60 s.	transverse <2%

The surface energy of the untreated BOPP film Radil T25 NT is less than 30 mN/m.

The surface energy of the polypropylene surface which is coated with adhesive composition is 52 mN/m after flame pretreatment and from 48 to 50 mN/m after corona treatment.

The opposing surface of the film, which is not coated with the adhesive composition, is provided with a reverse-side lacquer prior to coating by means of the customary technologies.

Example b: Flame pretreatment of the polypropylene films

Materials:

Film: Radil T 25 NT (BOPP film, RADICI), surface energy before flaming less than 30 mN/m

Fuel gas: propane (technical purity, LINDE)

Fuel gas/air mixture: approximate proportion = 1: 28 (v/v)

Technical parameters:

	Flaming side:	Outer side of the bale
	Web speed:	100 m/min
5	Burner/treatment roll distance:	3 mm
	Burner temperature:	38°C
	Temperature of the treatment roll:	15°C
	Flame temperature:	840°C
	Volume flow of mixture:	20 m³/h
10	Effective burner width:	32 cm

The film face flamed under this combination of parameters possesses a surface energy of 52 mN/m. A corona pretreatment suitable for comparison purposes normally achieves levels of from 48 to 50 mN/m.

Example c: Components of the adhesive compositions

- c1: Natural rubber TSR 5L obtainable from Weber&Schaer, Hamburg.
- c2: Hydrocarbon tackifier resin Hercotac 205 from Hercules B.V., Rijswijk, NL
- 20 c3: Ground chalk filler, Mikrosöhl 40, available from Vereinigte Kreidewerke Dammann KG, Söhlde
- c4: Plasticizer, paraffinic white oil, Shell Ondina G33 from Deutsche Shell AG, Hamburg
- c5: Ageing inhibitor, Lowinox ® 22M46 (2,2-methylenebis-[6-(1,1-dimethyl-ethyl)-4-methylphenol]) from GREAT LAKES
- 25 c6: Isocyanate crosslinker, 4,4'-diphenylmethane diisocyanate, product Suprasec DNR from ICI Polyurethanes, Löhne

Additionally for formulation of experiment B:

- 30 c7: UV crosslinking promoter, Ebecryl 140, dimerized trimethylol-propane tetraacrylate (C₂₄H₃₈O₉) from UCB Chemicals, Radcure Products, Drogenbos, Belgium

c8: Photoinitiator Irgacure 651, 2,2-dimethoxy-1,2-diphenylethan-1-one from Ciba Additive GmbH, Lampertheim, Germany

Example d: Adhesive composition formulations

5

The following formulations were used. All formulations are given in phr, i.e. based on 100 parts of rubber.

Formulation A:

10

Formulation for the chemical crosslinking of the adhesive composition based on a natural-rubber hot-melt pressure-sensitive adhesive by diisocyanate:

	Natural rubber, granulated	100
15	Hercotac 205	98
	Chalk	18
	White oil	4
	Ageing inhibitor	2
	Diisocyanate	0.9

20

Formulation B:

25 Formulation of the uncrosslinked adhesive composition based on a natural-rubber hot-melt pressure-sensitive adhesive:

	Natural rubber, granulated	100
	Hercotac 205	98
	Chalk	18
30	White oil	4
	Ageing inhibitor	2

Formulation C:

- 5 Formulation for the crosslinking by UV radiation of an adhesive composition based on a natural-rubber hot-melt pressure-sensitive adhesive:

	Natural rubber, granulated	100
	Hercotac 205	98
10	Chalk	18
	Ageing inhibitor	2
	Ebecryl 140	3.5
	Irgacure 651	4.5

15 **Example e: Preparation of the adhesive composition**

The adhesive compositions based on natural rubber were prepared without solvent in single-stage and multistage procedures.

20 **Example e1:**

In a first process step, a prebatch was prepared. The prebatch was prepared in accordance with the formulation designated VB-A in a Banbury compounder of type GK 1.4 N from Werner & Pfleiderer, Stuttgart.

25

Prebatch formulation VB-A

	Natural rubber, granulated	100
	Hercotac 205	18
	Chalk	18
	White oil	4
	Ageing inhibitor	2

Kneading chamber and rotors were conditioned at a temperature of 25°C, the rotary speed of the blades was 50 min⁻¹. The overall weight of the prebatch was 1.1 kg. All constituents were premixed in dry form and metered together with the white oil.

- 5 A mixing time of six minutes was sufficient to homogenize the constituents of the prebatch.

10 In a second process step, the natural-rubber hot-melt pressure-sensitive adhesive was mixed to completion. In a kneading machine of construction type LUK1.0 K3 from Werner & Pfleiderer, Stuttgart, all of the other additives were added for this purpose to the prebatch, so giving a natural-rubber hot-melt pressure-sensitive adhesive in accordance with the general formulation A. The prebatch was kneaded for ½ minute, then the entire tackifier resin in accordance with formulation F-A was added. The isocyanate was mixed in after 10 minutes.

15

Formulation F-A

Prebatch VB-A	142
Hercotac 205	80
Diisocyanate	0.9

20

The formulation F-A corresponds overall to the general formulation A of the natural-rubber hot-melt pressure-sensitive adhesive.

The overall weight of the hot-melt pressure-sensitive adhesive was 500 g. During the entire operation of mixing to completion, the chamber temperature was set at 80°C. The overall kneading time was 10 minutes.

25

Example e2

Example e1 was repeated.

30

To simplify emptying, the second step of the process, mixing to completion, was carried out using a kneading machine of the construction type VI U 20 L from Aachener Misch-

und Knetmaschinen-Fabrik Peter Küpper, Aachen, with a discharge screw. The prebatch was kneaded for ½ minute, then the entire tackifier resin in accordance with formulation F-A was added. The isocyanate was mixed in after 7 minutes.

- 5 The overall weight of the hot-melt pressure-sensitive adhesive was 12 kg. The chamber temperature was set at 80°C. The overall kneading time was 12 minutes, the emptying time 7 minutes.

- 10 The uncrosslinked formulation B and the UV-crosslinkable formulation C were prepared in analogy to Example e2 without the addition of isocyanate crosslinker to formulation B, and, respectively, by adding the UV crosslinker and the promoter in the case of formulation C.

15 **Example e3**

- 20 The natural-rubber hot-melt pressure-sensitive adhesive was prepared continuously with the aid of a twin-screw extruder L/D=36 from FARREL. In this case, the solid components, rubber and chalk, were supplied to the first conveying zone, the resin to the second conveying zone, and the liquid components were metered in.

Gravimetric metering devices, metering pumps and the screw speed were set so as to give a product rate of 10 kg/h with homogeneous mixing. The individual zones were temperature-controlled so as to give a product temperature of 75°C.

25 **Example e4**

The continuous preparation of the adhesive compositions based on natural-rubber hot-melt pressure-sensitive adhesives took place preferably by means of planetary roll extruders.

30

The planetary roll extruder was operated in each case with one, two or three roll cylinders, each roll cylinder being equipped with 6 planetary spindles and an approach ring having a free cross-section of 44 mm being used between each roll cylinder. The rotary speed of the central spindle was set at 100 revolutions/min. The maximum product

rate (Q_{\max}) up to which a homogenous compound was obtained was determined for each planetary roll extruder configuration.

In order to keep the natural-rubber granules meterable, they were treated with talc. The granules were produced using a cutting mill from Pallmann.

5

All of the components of the formulation were used to prepare, in a 50 kg powder mixture, a premix which was metered by way of a volumetric metering device into the filling section of the planetary roll extruder. The temperature-control circuits for the central spindle and the filling section (TC1 and TC2) were water cooled and each roll section was heated at 100°C. The table shows the maximum product rates achieved as a function of the number of roll cylinders.

10

Comparative Example	Number of roll cylinders	Q_{\max} [kg/h]
1	1	45
2	2	62
3	3	83

15

The components set out above in Example c were supplied continuously to the filling section of the planetary roll extruder, separately, by way of volumetric metering systems. The product exit temperature was measured as 122°C.

20

Example f: Coating of the adhesive composition

The continuously prepared and batchwise-prepared natural-rubber hot-melt pressure-sensitive adhesive with diisocyanate on the basis of formulation A, Example d, the non-crosslinked adhesive composition based on a natural-rubber hot-melt pressure-sensitive adhesive formulation B, Example d, and the UV crosslinker- and promoter-containing formulation C, Example d, were used for coating directly after the preparation process, using a flexurally rigid 2-roll applicator unit. For this purpose, a conveying extruder was used in order to convey the adhesive compositions.

25

An industry-standard 25 µm thick BOPP film, provided with an industry-standard primer coat based on isocyanate and an industry-standard release coat based on carbamate, was used and the adhesive compositions were applied with a film thickness of 18 µm.

- 5 The backing film described in a was coated directly, in a film thickness of 18 µm.

Coating took place on a pilot coating plant with a working width of 500 mm.

- 10 The coating of the film of adhesive onto the primed face of the film was carried out directly. A coating gap was established between the first coating roll and the second, web-carrying coating roll, in accordance with the application thickness. The first roll was heated at 100°C, the web-carrying roll cooled at 15°C. The adhesive composition emerging from the conveying extruder had a temperature of 120°C. Coating was carried out at 50 m/min. The properties of the adhesive tapes whose adhesive compositions were prepared by means of planetary roll extruders were investigated.

- 15 The properties were determined after storage for three days at 23°C

Example g: UV crosslinking of the adhesive composition

- 20 The specimens with formulation C were lined with a transparent siliconized polypropylene film and subsequently irradiated using a medium-pressure mercury lamp with an output of 105 W/cm at a speed of 40 m/min. The adhesive composition was irradiated on the one hand through the release film and on the other hand through the reverse of the backing.

Example h: Results

- 25 The test methods used are briefly characterized below:

The bond strength (peel strength) of the compositions was determined in accordance with AFERA 4001.

- 30 The shear strength of the adhesive compositions investigated was determined in accordance with PSTC 7 (Holding Power). All values stated were measured at room temperature and the stated load of 20 N with a bond area of 20x13 mm². The results are reported in minutes of holding time. The adhesion substrate used was steel.

35

In order to characterize the degree of crosslinking, the gel fraction in toluene was measured. For this purpose, square sections measuring 20 cm² were punched out and welded into a pouch made of a polyethylene spunbonded web (Tyvek from DuPont with a basis weight of approximately 55 g/m²). The specimens were then subjected to extraction with toluene for 3 days, with shaking. The toluene was changed each day. The toluene was subsequently replaced by hexane/heptane and the specimens were dried at 110°C. The gel was determined by differential weighing, taking into account extraction losses of the web and of the backing. The gel fraction is based on the rubber fraction in the formulation of the adhesive composition.

To determine the amount of composition applied, a circular specimen of known surface area is cut from the coated film and weighed. Subsequently, the adhesive composition is removed using petroleum spirit and the film, now free of adhesive composition, is reweighed. The amount of composition applied, in g/m², is calculated from the difference.

In order to characterize the tack with respect to cardboard, the adhesive tape is applied to the cardboard using a standard commercial manual roller and pressed on using a steel roller weighing 2 kg (overrolled twice). After a waiting time of 3 minutes, the adhesive tape is pulled off parallel at a speed of about 30 m/min and at an angle of approximately 130° to the cardboard surface. The tack with respect to cardboard is assessed qualitatively on the basis of the amount of paper fibres torn out, in comparison with a standard commercial adhesive packaging tape such as tesapack 4124 from Beiersdorf, whose tack is characterized as very good.

Property	Formulation A	Formulation B	Formulation C	
Crosslinking by	diisocyanate crosslinker	not crosslinked	UV crosslinking through the	
			release film	reverse
Adhesive composition applied [g/m ²]	18	18	18	18
Primer coat (standard)	no	no	yes	yes

Property	Formulation A	Formulation B	Formulation C	
Crosslinking by	diisocyanate crosslinker	not crosslinked	UV crosslinking through the	
			release film	reverse
Physical pretreatment of the film	flaming	flaming	corona before primer	corona before primer
Anchoring of composition	good	moderate	good	good
Gel fraction	-	0 %	72%	72%
Shear strength 20 N steel	>10,000 min	488	7200 min	>10,000 min
Bond strength on steel	3.1 N/cm	3.0 N/cm	1.8 N/cm	1.8 N/cm
Tack with respect to cardboard	good	good	good	good

002750 "440490

Claims

1. Adhesive tape comprising a backing comprising an oriented thermoplastic film and a coating comprising a solventlessly prepared pressure-sensitive adhesive composition based on non-thermoplastic elastomers such as natural rubber and tackifying resins, characterized in that the pressure-sensitive adhesive composition comprises a thermally labile crosslinking system.
2. Adhesive tape according to Claim 1, characterized in that the thermoplastic film comprises biaxially oriented HDPE, PVC or PET, monoaxially oriented polypropylene or biaxially oriented polypropylene.
3. Adhesive tape according to Claims 1 and 2, characterized in that the adhesive composition comprises a mixture comprising:
 - a) 100 parts by weight of natural rubber
 - b) 70-120 parts by weight of tackifying resins based on hydrocarbons
 - c) 5-30 parts by weight of fillers
 - d) 2-20 parts by weight of plasticizers
 - e) 0.1-15 parts by weight of a crosslinker system
 - f) 0.5-5 parts by weight of ageing inhibitors.
4. Adhesive tape according to one of Claims 1 to 3, characterized in that the crosslinker system is based on isocyanates, such as diisocyanates or polyisocyanates, and is used in particular at 0.1-5.0 parts by weight, especially 0.1-2.0 parts by weight.
5. Adhesive tape according to one of Claims 1 to 4, characterized in that the crosslinker system used comprises a mixture of at least one photoinitiator, used in particular at 0.1-5 parts by weight, and at least one polyfunctional (meth)acrylic ester, used in particular at 0.5-10 parts by weight.
6. Adhesive tape according to at least one of the preceding claims, characterized in that the pressure-sensitive adhesive composition is crosslinked by means of accelerated electrons or UV radiation.

7. Adhesive tape according to at least one of the preceding claims, characterized in that a coat of a primer is applied between the thermoplastic film and the adhesive layer.

8. Adhesive tape according to at least one of the preceding claims, characterized in that the thermoplastic film comprises a coating, for example a release coating.

9. Adhesive tape according to at least one of the preceding claims, characterized in that the adhesion of the adhesive composition to the thermoplastic film is improved by means of corona treatment or, preferably, flame pretreatment.

10. A process for producing an adhesive tape according to at least one of the preceding claims, characterized in that the pressure-sensitive adhesive composition based on non-thermoplastic elastomers is prepared continuously without solvent and without mastication in a continuously operating device having a filling section and a compounding section, the said process comprising

- a) feeding the solid components of the self-adhesive composition, such as elastomers and resins, into the filling section of the device, optionally feeding fillers, dyes and/or crosslinkers,
- b) transferring the solid components of the self-adhesive composition from the filling section to the compounding section,
- c) adding the liquid components of the self-adhesive composition, such as plasticizers, crosslinkers and/or further tackifying resins, to the compounding section,
- d) preparing a homogeneous self-adhesive composition in the compounding section, and
- e) discharging the self-adhesive composition.

11. Use of the adhesive tape according to one of Claims 1 to 10, with a backing based on biaxially oriented polyolefins, on plasticized PVC or PET, as an adhesive packaging tape with good carton sealing and with good tack with respect to paper, especially recycled paper, sufficient cohesion, good packaging security, and easy unrolling.

12. Use of the adhesive tape according to one of Claims 1 to 11, with a backing based on monoaxially oriented polyolefins, as an adhesive packaging tape, strapping tape, for bundling and palletizing cardboard packaging and other goods.

Abstract

Adhesive tape comprising a backing comprising an oriented thermoplastic film and a coating comprising a solventlessly prepared pressure-sensitive adhesive composition based on non-thermoplastic elastomers such as natural rubber and tackifying resins, characterized in that the pressure-sensitive adhesive composition comprises a thermally labile crosslinking system.

05641014-081700

COMBINATION DECLARATION & POWER OF ATTORNEY

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled „**Adhesive packaging tape with natural-rubber hot-melt pressure sensitive adhesive**“
the specification of which is attached hereto.

-OR-

was filed on _____ as

Application Serial No. _____ and was amended _____

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)			Priority Claimed
<u>199 39 076.2</u> (Number)	<u>Germany</u> (Country)	<u>18/08/1999</u> (Day/Month/Yr. Filed)	[X] yes [] no
_____	_____	_____	[X] yes [] no
(Number)	(Country)	(Day/Month/Yr. Filed)	

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

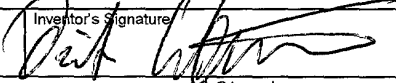

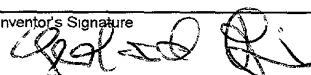

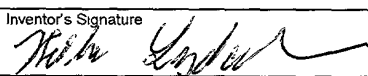
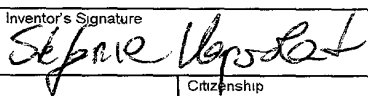
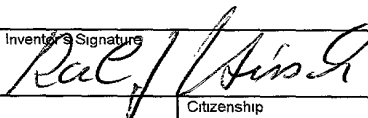
_____	_____	_____
(Application Serial No.)	(Filing Date)	(Status)
		(patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punished by fine or imprisonment, or both under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

POWER OF ATTORNEY: As a named Inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith:


Kurt G. Briscoe, Reg. No. 33,141; William C. Gerstenzang, Reg. No. 27,552; Carmella A. O'Gorman, Reg. No. 33,749; and Stephen G. Ryan, Reg. No. 39,015 all of 660 White Plains Road, Tarrytown, New York 10591-5144; William R. Robinson, Reg. No. 27,224; Mark A. Montana, Reg. No. 44,948 all of 721 Route 202-206, Bridgewater, New Jersey 08807; Lorimer P. Brooks, Reg. No. 15,155; Davy E. Zoneraich, Reg. No. 37,267 all of 805 Third Avenue, 9th Floor, New York, NY 10022, my attorneys with full power of substitution and revocation

Send Correspondence To: Norris McLaughlin & Marcus, P.A. 660 White Plains Road Tarrytown, N. Y. 10591-5144	Direct Telephone Calls To: (914) 332-1700
--	---

Full Name Of Sole or First Inventor Dr. Dieter Wenninger	Inventor's Signature 	Date 27.06.00
Residence Keplerstraße 12, D-22765 Hamburg, Germany	Citizenship German	
Post Office Address Keplerstraße 12, D-22765 Hamburg, Germany		
Full Name Of Second Inventor Dr. Werner Karmann	Inventor's Signature 	Date 29.6.00
Residence Stolpmünder Straße 11, D-22147 Hamburg, Germany	Citizenship German	
Post Office Address Stolpmünder Straße 11, D-22147 Hamburg, Germany		
Full Name Of Third Inventor Dr. Gerhard Bir	Inventor's Signature 	Date 28.06.00
Residence Barckhusendamm 2, D-22117 Hamburg, Germany	Citizenship German	
Post Office Address Barckhusendamm 2, D-22117 Hamburg, Germany		
Full Name Of Fourth Inventor Sven Hansen	Inventor's Signature 	Date 28.06.00
Residence Lutterothstraße 23, D-20255 Hamburg, Germany	Citizenship German	
Post Office Address Lutterothstraße 23, D-20255 Hamburg, Germany		
Full Name Of Fifth Inventor Heiko Leydecker	Inventor's Signature 	Date 28.06.00
Residence Kreienredder 100, D-24730 Neustadt/Holstein, Germany	Citizenship German	
Post Office Address Kreienredder 100, D-24730 Neustadt/Holstein, Germany		
Full Name Of Sixth Inventor Stefanie Kaprolat	Inventor's Signature 	Date 03.07.00
Residence Hintergasse 52, D-35216 Biedenkopf, Germany	Citizenship German	
Post Office Address Hintergasse 52, D-35216 Biedenkopf, Germany		
Full Name Of Seventh Inventor Ralf Hirsch	Inventor's Signature 	Date 4.7.00
Residence Finkenweg 2, D-25451 Quickborn, Germany	Citizenship German	
Post Office Address Finkenweg 2, D-25451 Quickborn, Germany		

0043043700

0024304403430

Full Name Of Eighth Inventor Klaus Massow		Inventor's Signature 	Date 08/07/10
Residence Stolpmünder Straße 35D, D-22147 Hamburg, Germany		Citizenship German	
Post Office Address Stolpmünder Straße 35D, D-22147 Hamburg, Germany			
Full Name Of Ninth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Tenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Eleventh Inventor		Inventor's Signature	Date
Residence,		Citizenship	
Post Office Address			
Full Name Of Twelfth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Thirteenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Fourteenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Fifteenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			
Full Name Of Sixteenth Inventor		Inventor's Signature	Date
Residence		Citizenship	
Post Office Address			